Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A pharmaceuticals characterized by general formula (I)

$$Z\text{-}L\text{-}V \quad (I)$$

wherein

V denotes a peptide with a binding sequence $-X^1-X^2$-Val-Tyr-Ile-His-Pro-$X^8-X^9-X^{10}$, (SEQ ID NO. 1)

L denotes bond or a linker,

Z denotes a chelating agent suitable for ~~group that optionally can~~ carrying a radionuclide ~~an imaging moiety~~ M,

X^1 denotes-NY_1-$(CH_2)_m$-CO- where m is an integer from 1 to 10 and Y_1 is H or an alkyl- or aryl-containing substituent,

X^2 denotes Arg, N-alkylated Arg, or ~~a Arg mimetic~~ Phe[4-guanidino] or Gly-4-piperidyl[N-amidino],

X^8 denotes Gly, Phe, Phg, Hph, Bip, Ala, Tyr, His, Trp or Nal, ~~SEQ ID NO. 1~~

X^9 and X^{10} denote, independent of each other, Pro, Arg, His, Ala, Phe, Glu, Leu, Val, Ile, Met, Trp, Asp or Lys ~~SEQ ID NO. 1~~ and where X^8, X^9 and X^{10} together constitute an ACE cleavage site;

and wherein the residues Val and Ile at position 3 and 5 respectively may optionally be replaced with the amino acids cysteine or homocysteine capable of forming a bridging unit wherein the bridge contain~~ings~~ a $-CH_2-CH_2-$, $-S-CH_2-$, $-S-CH_2-S-$, lactam or –-S-S- unit,

Z forms a bond with the amino acid X^1 optionally through the linker L, and

M ~~where present denotes an imageable moiety capable of detection either directly or indirectly in a diagnostic imaging procedure~~ is selected from ^{67}Ga, ^{111}In, ^{81m}Kr, ^{99}Mo, ^{98m}Tc, ^{201}Tl, ^{68}Ga and ^{82}Rb.

2. (Currently Amended) A pharmaceutical according to claim 1 wherein the amino acid of X^1 , X^2, X^8, X^9, X^{10} are independently selected from

X^1 denoting Gly

X^2 denoting Arg or N-Methyl-Arg

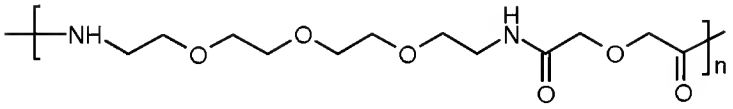
X^8 denoting Phe

X^9 denoting Pro, Arg, His, Ala, Phe, Glu, Leu, Val, Ile, Met, Trp, Asp or Lys ~~SEQ ID NO. 1~~ and

X^{10} denoting Pro, Arg, His, Ala, Phe, Glu, Leu, Val, Ile, Met, Trp, Asp or Lys ~~SEQ ID NO. 1~~.

3. (Currently Amended) A pharmaceutical according to claim 1 further comprising one or more biomodifier groups ~~are attached to any positions of the V and L groups of formula (I)~~

of formula (IV)

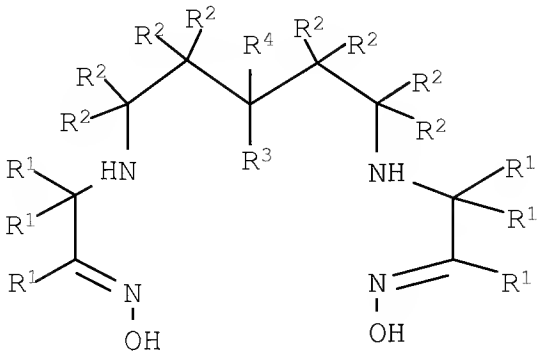


(IV)

wherein n equals an integer from 1 to 10 and where the C-terminal unit forms an amide bond attached to any positions of the V and L groups of formula (I).

4. (Cancelled) A pharmaceutical according to claim 1 wherein Z denotes a chelating agent.

5. (Original) A pharmaceutical according to claim 4 wherein Z denotes the chelating agent of formula (VII)



(VII)

wherein:

each R^1, R^2, R^3 and R^4 is independently H or C_{1-10} alkyl, C_{3-10} alkylaryl, C_{2-10} alkoxyalkyl, C_{1-10} hydroxyalkyl, C_{1-10} alkylamine, C_{1-10} fluoroalkyl, or 2 or more R groups, together with the atoms to which they are attached form a carbocyclic, heterocyclic, saturated or unsaturated ring.

6. (Cancelled) A pharmaceutical according to claim 5 wherein M represents an imageable moiety for the use in diagnosis particularly in *in vivo* diagnosis comprising a moiety which emit or cause to emit detectable radiation, a moiety which affect local electromagnetic fields, moieties which absorb or scatter radiation energy, heavy metals and compounds thereof and moieties which generate a detectable substance.

7. (Currently Amended) A pharmaceutical according to claim 5~~6~~ wherein M represents a gamma emitting moiety for Radio or SPECT imaging ~~comprising~~ selected from ^{67}Ga, ^{111}In, ~~^{123}I, ^{125}I, ^{131}I~~, ^{81m}Kr, ^{99}Mo, ^{99m}Tc and ^{201}Tl ~~and ^{133}Xe.~~

8. (Currently Amended) A pharmaceutical according to claim 5~~6~~ wherein M represents a radio emitter with positron emitting properties for PET imaging comprising ~~^{11}C, ^{18}F,~~ ^{68}Ga, ~~^{13}N, ^{15}O~~ and ^{82}Rb.

9. (Currently Amended) A pharmaceuticals according to claim 2~~1~~ characterized by ~~general formula (I)~~

~~Z-L-V-~~ ~~(I)~~

~~wherein~~

~~V denotes a peptide with a binding sequence X^1-X^2-Val-Tyr-Ile-His-Pro-X^8-X^9-X^{10}, SEQ ID NO. 1 wherein the amino acid of X^1, X^2, X^8, X^9, X^{10} are independently selected from~~

~~X^1 denoting Gly~~

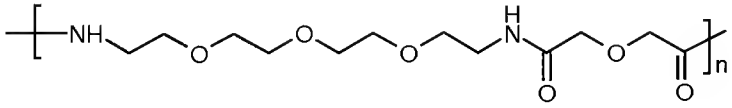
~~X^2 denoting Arg or N-Methyl-Arg~~

~~X^8 denoting Phe~~

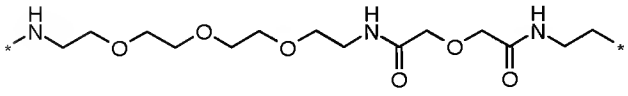
~~X^9 denoting Pro, Arg, His, Ala, Phe, Glu, Leu, Val, Ile, Met, Trp, Asp or Lys SEQ ID NO. 1 and~~

~~X^{10} denoting Pro, Arg, His, Ala, Phe, Glu, Leu, Val, Ile, Met, Trp, Asp or Lys SEQ ID NO. 1.~~

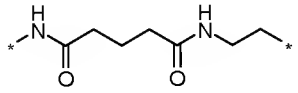
wherein L denotes a bond or a linker selected from compounds of formula NH-(CH2)m- optionally combined with –CO-(CH2)m-CO- where m denotes a positive integer from 1 to 10, one or more units of compounds of formula (IV) wherein n is an integer from 1 to 10, compounds of formula (X) or (VI)



Formula (IV)

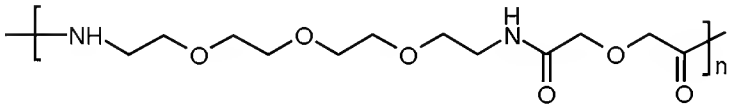


Formula (X)



Formula (VI)

~~Z denotes a chelating agent of formula (VII) that optionally can carry an imaging moiety M, and one or more biomodifier groups selected from monodisperse PEG building block comprising 1 to 10 units of said building block or the compound of formula IV,~~



~~Formula (IV)~~

~~wherein n equals an integer from 1 to 10 are attached to any positions of the V and L groups of formula (I).~~

10. (Original) Pharmaceutical formulation comprising a pharmaceutical of formula (I) of claim 1 together with one or more pharmaceutical acceptable additives and/or excipients.

11. (Cancelled) A kit for the preparation of a radiopharmaceutical composition of formula (I) comprising a peptide-chelate conjugate and a reducing agent.